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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.		
09/970,641	10/03/2001	Chin-Shiou Huang	ASPI-002/03US	2745		
75	90 05/01/2003			•		
PENNIE & EDMONDS LLP 1155 Avenue of the Americas			EXAMINER FRIEND, TOMAS H F			
			ART UNIT	PAPER NUMBER		
			1639			
			DATE MAILED: 05/01/2003	7		

Please find below and/or attached an Office communication concerning this application or proceeding.

		Application I	No.	Applicant(s)		
		09/970,641	_	HUANG, CHIN-SHIOU		
	Office Action Summary	Examiner		Art Unit		
		Tomas Friend	d	1639		
	The MAILING DATE of this communication appears on the cover sheet with the correspondence address Period for Reply					
A SHORTENED STATUTORY PERIOD FOR REPLY IS SET TO EXPIRE 1 MONTH(S) FROM THE MAILING DATE OF THIS COMMUNICATION. Extensions of time may be available under the provisions of 37 CFR 1.136(a). In no event, however, may a reply be timely filed after SIX (6) MONTHS from the mailing date of this communication. If the period for reply specified above is less than thirty (30) days, a reply within the statutory minimum of thirty (30) days will be considered timely. If NO period for reply is specified above, the maximum statutory period will apply and will expire SIX (6) MONTHS from the mailing date of this communication. Failure to reply within the set or extended period for reply will, by statute, cause the application to become ABANDONED (35 U.S.C. § 133). Any reply received by the Office later than three months after the mailing date of this communication, even if timely filed, may reduce any earned patent term adjustment. See 37 CFR 1.704(b).						
Status 1)⊠	Responsive to communication(s) filed on 12 A	Juguet 2002				
2a)□		is action is no	n-final			
3)□	<i>,</i> —			secution as to the i	merits is	
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213. Disposition of Claims						
·	Claim(s) <u>1-27</u> is/are pending in the application.					
•	4a) Of the above claim(s) is/are withdraw		deration.			
	Claim(s) is/are allowed.					
	Claim(s) is/are rejected.					
	Claim(s) is/are objected to.					
8)⊠	Claim(s) <u>1-27</u> are subject to restriction and/or e	election require	ement.			
Applicati	on Papers	·				
9)[The specification is objected to by the Examiner	r.				
10)[] 7	Γhe drawing(s) filed on is/are: a)□ accep	oted or b) 🔲 obj	ected to by the Exan	niner.		
	Applicant may not request that any objection to the		· ·			
11) 🔲 🗆	The proposed drawing correction filed on	_is: a)∏ appr	oved b)⊡ disapprov	ved by the Examiner.		
_	If approved, corrected drawings are required in rep	•	action.			
12) The oath or declaration is objected to by the Examiner.						
Priority under 35 U.S.C. §§ 119 and 120						
13) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f).						
a) ☐ All b) ☐ Some * c) ☐ None of:						
	1. Certified copies of the priority documents					
	2. Certified copies of the priority documents have been received in Application No					
 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 						
14) Acknowledgment is made of a claim for domestic priority under 35 U.S.C: § 119(e) (to a provisional application).						
a) ☐ The translation of the foreign language pro	visional applic	cation has been rece	eived.	,	
15) ☐ Acknowledgment is made of a claim for domestic priority under 35 U.S.C. §§ 120 and/or 121. Attachment(s)						
1) Notice	e of References Cited (PTO-892) e of Draftsperson's Patent Drawing Review (PTO-948) nation Disclosure Statement(s) (PTO-1449) Paper No(s)	5)	Notice of Informal P	(PTO-413) Paper No(s). atent Application (PTO-1		

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Detailed Action

Status of the Application

Receipt is acknowledged of a preliminary amendment on 03 October 201 and an information disclosure statement on 12 August 2002 (Paper Nos. 5 and 6).

Status of the Claims

Claims 1-27 are pending in the present application and are subject to restriction and election of species requirements.

Sequence Compliance

This application contains sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 CFR 1.821 (a) (1) and (a) (2). See, for example, page 32 of the specification. However, this application does not comply with the requirements of 37 CFR 1.821 through 1.825 for the reason(s) set forth on the attached Notice to Comply with Requirements for Patent Applications Containing Nucleotide Sequence and/or Amino Acid Sequence Disclosures. Applicant must comply with the requirements of the sequence rules (37 CFR 1.821 - 1.825) before the application can be further examined under 35 U.S.C. §§ 131 and 132.

Applicant's response to this office action must bring this application into compliance with the sequence rules. Applicant is advised to inspect the specification, claims, and figures to identify and bring into compliance all sequence disclosures that are encompassed by the definitions for nucleotide and/or amino acid sequences set forth in 37 CFR 1.821 (a) (1) and (a) (2). Please see the attached Notice to Comply with the Sequence Rules for further information.

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Restriction

1. Restriction to one of the following inventions is required under 35 U.S.C. 121:

- I. Claims 1-19, drawn to a surface imprint composition comprising a matrix material defining imprint cavities of a template molecule wherein a substantial fraction of the imprint cavities are localized at or near the surface of the matrix material, classified in class 521, subclass 99.
- II. Claims 20-26, drawn to a plurality of surface imprint compositions according to claim 1, classified in class 435, and one of digests 22-42, depending on the compositions of the matrix material, template molecule, and the presence or absence of the template molecule.
- III. Claim 27, drawn to a surface imprint composition comprising a matrix material defining imprint cavities of a template molecule wherein a substantial fraction of the imprint cavities are oriented, classified in class 521, subclass 99.

The inventions are distinct, each from the other because:

- 2. Inventions I-III are different and patentably distinct compositions. For example, Inventions I and III are each a single imprint composition defining imprint cavities of a single template while Invention II is a plurality of surface imprint compositions that, together, define imprint cavities of a plurality of different imprint cavities of different templates. Inventions I and III are patentably distinct compositions because Invention I does not require imprint cavities to be oriented as required in Invention III and Invention III does not require a substantial fraction of imprint cavities to be at or near the surface as required in Invention I.
- 3. Because these inventions are distinct for the reasons given above and
 - a. have acquired a separate status in the art as shown by their different classification;
 - b. have different and separately burdensome: manual and/or computer: structure, name and bibliographical searches; and
 - c. have divergent subject matter, restriction for examination purposes as indicated is proper.

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4. Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143)

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a diligently filed petition under 37 CFR 1.48(b) and by the fee required under CFR 1.17(h).

Election of Species

- 5. This application contains claims directed to patentably distinct species of the claimed invention. IN addition to electing one of Inventions I or II, applicant is required a species for each of the following A-C.
 - A. species of matrix material (e.g. from the list in claim 3) including identity of heat sensitive compound, if present,
 - B. number and ultimate species of template molecule(s) (identified by molecule names or structures but NOT as a molecular class such as proteins, nucleic acids, or small molecules) AND
 - C. what part(s) of the template molecule(s) correspond to the imprint cavities.
- 6. The species are distinct, each from the other, because they have different structures with different chemical, physical, and/or pharmacological properties. Therefore, different issues of enablement and patentability apply to each species and each species represents patentably distinct subject matter.
- 7. Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claims 1, 20, and 27 are generic.

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Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

8. Any inquiry concerning this communication or earlier communications from the examiner should be directed to **Tomas Friend**, telephone number (703) 308-4548. The examiner's schedule is normally four, ten-hour days per week that includes Saturdays.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Andrew Wang can be reached on (703) 306-3217. The fax phone number for the organization where this application or proceeding is assigned is (703) 308-2742.

Any inquiry of a general nature or relating to the status of this application or proceeding should be directed to the receptionist at (703) 308-1235.

Tomas Friend, Ph.D. 26 April 2003

ANDREW WANG SUPERVISORY PATENT EXAMINER TECHNOLOGY CENTER 1600

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NOTICE TO COMPLY WITH REQUIREMENTS FOR PATENT APPLICATIONS CONTAINING NUCLEOTIDE SEQUENCE AND/OR AMINO ACID SEQUENCE DISCLOSURES

Applicant must file the items indicated below within the time period set the Office action to which the Notice is attached to avoid abandonment under 35 U.S.C. § 133 (extensions of time may be obtained under the provisions of 37 CFR 1.136(a)).

The nucleotide and/or amino acid sequence disclosure contained in this application does not comply with the requirements for such a disclosure as set forth in 37 C.F.R. 1.821 - 1.825 for the following reason(s):

X	1. This application clearly fails to comply with the requirements of 37 C.F.R. 1.821-1.825. Applicant's attention is directed to the final rulemaking notice published at 55 FR 18230 (May 1, 1990), and 1114 OG 29 (May 15, 1990). If the effective filing date is on or after July 1, 1998, see the final rulemaking notice published at 63 FR 29620 (June 1, 1998) and 1211 OG 82 (June 23, 1998).
X	2. This application does not contain, as a separate part of the disclosure on paper copy, a "Sequence Listing" as required by 37 C.F.R. 1.821(c).
X	3. A copy of the "Sequence Listing" in computer readable form has not been submitted as required by 37 C.F.R. 1.821(e).
	4. A copy of the "Sequence Listing" in computer readable form has been submitted. However, the content of the computer readable form does not comply with the requirements of 37 C.F.R. 1.822 and/or 1.823, as indicated on the attached copy of the marked -up "Raw Sequence Listing."
	5. The computer readable form that has been filed with this application has been found to be damaged and/or unreadable as indicated on the attached CRF Diskette Problem Report. A Substitute computer readable form must be submitted as required by 37 C.F.R. 1.825(d).
	6. The paper copy of the "Sequence Listing" is not the same as the computer readable from of the "Sequence Listing" as required by 37 C.F.R. 1.821(e).
	7. Other:
Αp	plicant Must Provide:
X	An initial or substitute computer readable form (CRF) copy of the "Sequence Listing".
X	An initial or <u>substitute</u> paper copy of the "Sequence Listing", as well as an amendment directing its entrinto the specification.
X	A statement that the content of the paper and computer readable copies are the same and, where applicable, include no new matter, as required by 37 C.F.R. 1.821(e) or 1.821(f) or 1.821(g) or 1.825(b) or 1.825(d).
Fo	questions regarding compliance to these requirements, please contact:
Fo	Rules Interpretation, call (703) 308-4216 CRF Submission Help, call (703) 308-4212 tentIn Software Program Support Technical Assistance

PLEASE RETURN A COPY OF THIS NOTICE WITH YOUR REPLY